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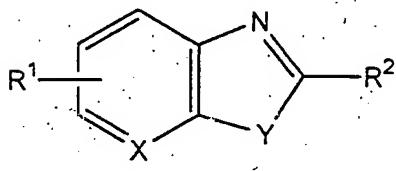
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LISTING OF CLAIMS:

-- Claim 1. (Original)

A method for inhibiting 5-lipoxygenase in a subject, comprising administering a compound of formula (I) or a pharmaceutically acceptable salt thereof to the subject in an amount effective for the inhibition of 5-lipoxygenase:



(I)

wherein

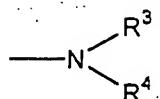
X is CH or N;

Y is S or O;

R¹ is H, OH, halogen, C₁₋₆ alkyl, nitro, cyano, amino, di-C₁₋₆ alkylamino, C₁₋₆ alkoxy, C₁₋₆ hydroxyalkyl or C₁₋₆ alkylcarbonyl; and

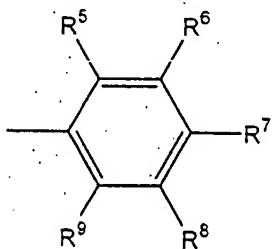
R² is

(i)



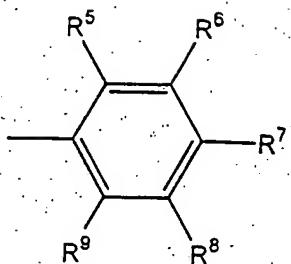
wherein R³ is H or C₁₋₆ alkyl;

R⁴ is



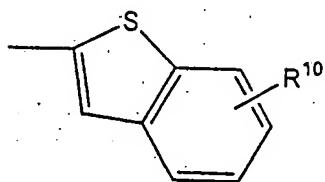
wherein R⁵, R⁶, R⁷, R⁸ and R⁹ are independently H, OH, halogen, C₁₋₆ alkyl, C₁₋₆ haloalkyl, nitro, cyano, amino, di-C₁₋₆ alkylamino, mercapto, C₁₋₆ mercaptoalkyl, halogen-substituted C₁₋₆ mercaptoalkyl, phenylazo, C₁₋₆ alkylphenylazo, C₁₋₆ alkylcarbonyl, C₁₋₆ alkoxy or C₁₋₆ hydroxyalkyl,

(ii)



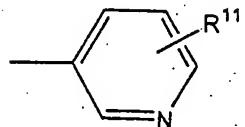
wherein R⁵, R⁶, R⁷, R⁸ and R⁹ are as defined in (i),

(iii)



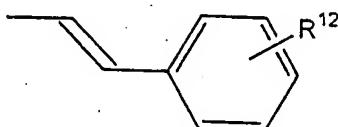
wherein R¹⁰ is H or C₁₋₆ alkyl,

(iv)



wherein R¹¹ is H, C₁₋₆ alkyl, halogen, mercapto or C₁₋₆ mercaptoalkyl, or

(v)



wherein R¹² is H, OH, halogen, C₁₋₆ alkyl, nitro, cyano, amino, di-C₁₋₆ alkylamino, C₁₋₆ alkylcarbonyl, C₁₋₆ alkoxy or C₁₋₆ hydroxyalkyl.

-- Claim 2. (Original)

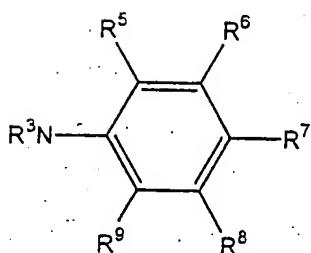
The method of claim 1, which is used for preventing or treating a leukotriene-related disease selected from the group consisting of: asthma, pertussis, psoriasis, rheumatic arthritis, arthritis, inflammatory bowel disease, cystic fibrosis, acute/chronic bronchitis, gout, sepsis, cardiac myoischemia, cardiac anaphylaxis, cerebrovascular convulsion, ischemia and allergic rhinitis.

-- Claim 3. (Original)

The method of claim 2, wherein the disease is asthma.

-- Claim 4. (Original)

The method of claim 1, wherein R² is



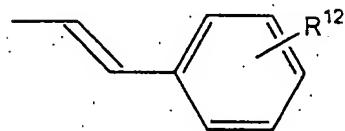
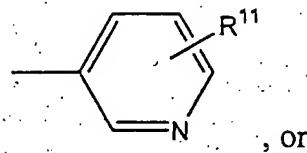
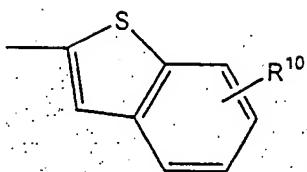
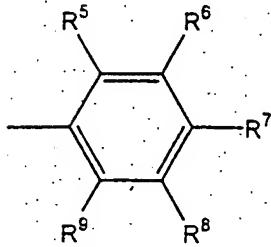
wherein R³, R⁵, R⁶, R⁷, R⁸ and R⁹ are as defined in claim 1.

-- Claim 5. (Original)

The method of claim 4, wherein R¹ is H, halogen, C₁₋₆ alkyl or nitro; and R⁵, R⁶, R⁷, R⁸ and R⁹ are independently H, halogen, C₁₋₆ alkyl or phenylazo.

-- Claim 6. (Original)

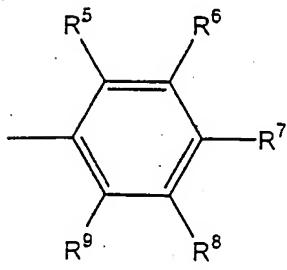
The method of claim 1, wherein R² is

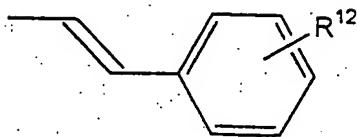
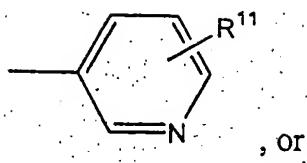


wherein R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, R¹¹ and R¹² are as defined in claim 1.

-- Claim 7. (Original)

The method of claim 6, wherein R¹ is H or C₁₋₆ alkyl; and R² is





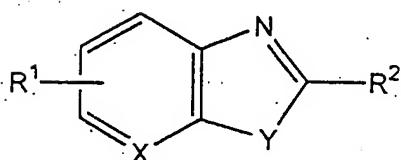
wherein R⁵, R⁶, R⁷, R⁸ and R⁹ are, independently, H, halogen, C₁₋₆ alkyl, C₁₋₆ haloalkyl, nitro, cyano, amino, di-C₁₋₆ alkylamino, mercapto, C₁₋₆ mercaptoalkyl, halogen-substituted C₁₋₆ mercaptoalkyl or C₁₋₆ alkoxy;

R^{¹¹} is as defined in claim 1; and

R^{¹²} is H, halogen or C₁₋₆ alkyl.

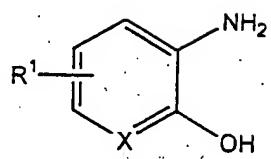
-- Claim 8. (Amended)

A method for preparing a compound of formula (I)

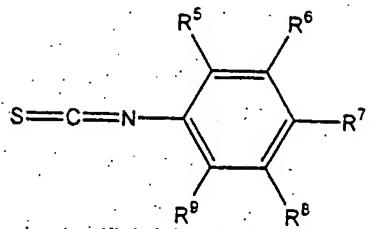


(I)

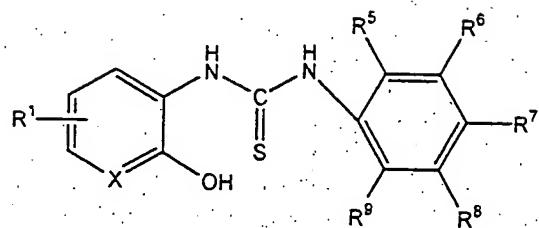
comprising the steps of: (a) reacting a compound of formula (II) with a compound of formula (III) in an organic solvent to synthesize a thiourea intermediate of formula (IV); and (b) reacting the thiourea intermediate of formula (IV) with an acid to obtain a compound of formula (Ia) or (Ib):



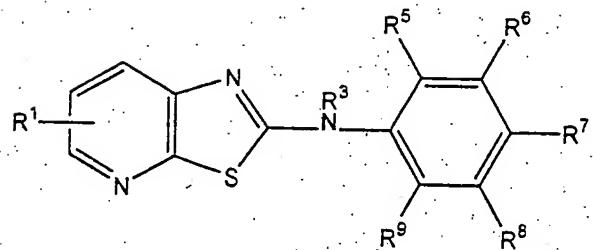
(II)



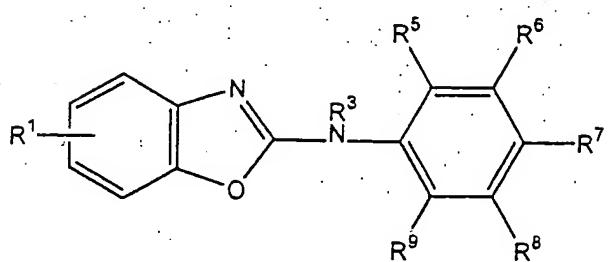
(III)



(IV)



(Ia)



(Ib)

wherein R₁, R₃, R₅, R₆, R₇, R₈ and R₉ are as defined in claim 1.

R¹ is H, OH, halogen, C₁₋₆ alkyl, nitro, cyano, amino, di-C₁₋₆ alkylamino, C₁₋₆ alkoxy, C₁₋₆ hydroxyalkyl or C₁₋₆ alkylcarbonyl; and
wherein R³ is H or C₁₋₆ alkyl;
wherein R⁵, R⁶, R⁷, R⁸ and R⁹ are independently H, OH, halogen, C₁₋₆ alkyl, C₁₋₆ haloalkyl, nitro, cyano, amino, di-C₁₋₆ alkylamino, mercapto, C₁₋₆ mercaptoalkyl, halogen-substituted C₁₋₆ mercaptoalkyl, phenylazo, C₁₋₆ alkylphenylazo, C₁₋₆ alkylcarbonyl, C₁₋₆ alkoxy or C₁₋₆ hydroxyalkyl.

-- Claim 9. (Original)

The method of claim 8, wherein the acid in step (b) is selected from the group consisting of trifluoroacetic acid, phosphoric acid, sulfuric acid, hydrochloric acid and nitric acid.